## CLAIM AMENDMENTS

- 1. (Currently Amended) A method of identifying a nucleic acid in a sample, comprising:
  - a) combining the sample with a polynucleotide probe comprising-a-ecquence-identical or complementary to at-least-10 consecutive nucleotides contained in SEQ I/D NO:224, such that the probe hybridizes specifically to the nucleic acid if the nucleic acid encodes human telomerase reverse transcriptase (hTRT) or fragment thereof;
    - b) detecting any hybrid formed as a result of a); and
  - c) identifying the nucleic acid as encoding at least a portion of human telemerase reverse transcriptase (hTRT) hTRT or tragment thereof if the hybrid is detected;

wherein the probe hybridizes specifically to a DNA having the sequence of the hTRT encoding region of SEO. ID NO:224 at 5°C to 25°C below T<sub>m</sub> in equeous solution at 1 M NaCl;

wherein T<sub>m</sub> is the melting temperature of double-stranded DNA having the sequence of said encoding region under the same reaction conditions.

- (Currently Amended) A method of detecting a nucleic acid that encodes hTRT or fragment thereof in a sample, comprising:
  - a) combining the sample with a polynucleotide probe such that the probe hybridizes specifically to a nucleic acid comprising at least 100 consecutive nucleotides contained in SEQ ID NO:224 if present in the sample; and
    - b) detecting any hybrid formed as a result of a);

wherein the polynucleotide probe comprises consists essentially of a sequence identical or complementary to at-least 25 or more consecutive nucleotides contained in from the hTRT encoding region of SEQ ID NO:224.

- 3. (Original) The method of claim 2, wherein the hTRT nucleic acid is human genomic DNA.
- (Currently Amended) The method of claim 2, wherein the hTRT nucleic acid is human-mRNA mRNA or cDNA.
- (Currently Amended) The method of claim 2, wherein the hTRT nucleic acid comprises at least 250 or more nucleotides of SEQ ID NO:224.
- (Currently Amended) The method of claim 2, wherein the hTRT nucleic acid comprises at least 500 or more nucleotides of SEQ ID NO:224.

- (Currently Amended) The method of claim 2, wherein the probe comprises a sequence identical
  or complementary to at-least-30 or more consecutive nucleotides contained in from the hTRT
  encoding region of SEQ ID NO:224.
- (Currently Amended) The method of claim 2, wherein the probe comprises a sequence identical
  or complementary to at-least 50 or more consecutive nucleotides contained in from the hTRT
  encoding region of SEQ ID NO:224.
- (Currently Amended) The method of claim 2, wherein the probe comprises a sequence identical
  or complementary to at-least 100 or more consecutive nucleotides contained-in from the hTRT
  encoding region of SEQ ID NO:224.
- (Original) The method of claim 2, wherein the probe comprises a sequence not contained in SEO. ID NO:62.
- (Original) The method of claim 9, wherein the probe comprises a sequence not contained in SEQ. ID NO:62.
- 12. (Original) The method of claim 2, wherein the sample is a human biological sample.
- 13. (Currently Amended) A method of identifying a nucleic acid in a sample, comprising:
  - a) combining the sample with a polynucleotide primer containing-a sequence-identical-er complementary to at least 10-censessuive nucleotides-contained in SEQ-10 Not224, under conditions that the primer amplifies specifically primes amplification of the nucleic acid if the nucleic acid encodes human telomorase reverse transcriptase (hTRT) or fragment thereof;
    - b) detecting any amplification product formed as a result of a); and
  - c) identifying the nucleic acid as encoding at least-a-portion of hTRT or fragment thereof if
    the amplification product is detected;
    - wherein the primer hybridizes specifically to a DNA having the sequence of the hTRT encoding region of SEQ. ID NO:224 at 5°C to 25°C below T<sub>n</sub> in squeous solution at 1.M Machine
  - wherein  $\mathbf{I}_m$  is the melting temperature of double-stranded DNA having the sequence of said encoding region under the same reaction conditions.

- (Currently Amended) A method of detecting a nucleic acid encoding at least a portion of hTRT gr fragment thereof in a sample, comprising:
  - a) combining the sample with a-polymeteotide-primer-such-that the primer amplifies
     <u>polymeteotide primers so as to prime amplification of</u> nucleic acid encoding <del>at least a pertion of</del>
     <u>hTRT or fragment thereof if present in the sample;</u> and
    - b) detecting any amplified product formed as a result of a);
  - wherein the pelywoleotide primes—comprises gach of said primers consists assentially of a sequence identical or complementary to at-least 15 or more consecutive nucleotides centained in from the THT encoding region of SECI ID NO:224.
- 15. (Currently Amended) The method of claim 14, wherein the polyeucleokide-primer-comprises sach of said primers consists essentially of a sequence identical or complementary to at-least 30 gr more consecutive nucleotides centained in from the hTRT encoding region of SEQ ID NO:224.
- (Currently Amended) The method of claim 14, wherein the-polynucleotide privace comprises each
  of said primers consists assantially of a sequence identical or complementary to at-least 50 or
  more consecutive nucleotides centained-in from the hTRT encoding region of SEQ ID NO:224.
- 17. (Original) The method of claim 14, wherein the sample is a human biological sample.
- 18. (Original) The method of claim 14, wherein the sample comprises human genomic DNA.
- (Currently Amended) The method of claim 14, wherein the sample comprises human-mRNA hTRT mRNA or cDNA.
- 20. CANCELLED.
- (Original) The method of claim 14, wherein the primer-comprises primers comprise a sequence not contained in SEQ, ID NO:62.
- 22. GANCELLED

- 23. (Withdrawn) (Currently Amended) A combination of oligonucleotide primers for PCR amplification for use in detecting an hTRT nucleic soid according to daim 14, comprising a first primer. that hybridizes to a polyneolootide-conclusing of SEC 45-NO:224 under-stringent amplification conditione, and a second primer that hybridizes to the complement of said nucleic acid under-stringent amplification-conditione wherein each primer consists essentially of a sequence identical or complementary to 15 or more consecutive nucleotides from the hTRT encoding region of SECID NO:224.
- (Withdrawn) The combination of primers of claim 23, wherein either each primer eemprises between consists of 15-30 nucleotides.
- (Withdrawn) The combination of primers of claim 23, wherein either each primer comprises between consists of 20-25 nucleotides.
- (Withdrawn) The combination of primers of claim 23, wherein 50% or more of the nucleotides of either each primer are guanine and/or cytosine.
- (Withdrawn) (Currently Amended) A PCR product that hybridizee-under-stringent-conditions-to-a
  polynucleotide having a sequence-condicting-of formed while undertaking the detection method of
  claim 14, comprising 15 or more contiguous nucleotides of the hTRT encoding region of SEQ ID
  NO:224 or its complement.
- (Withdrawn) (Currently Amended) A hybridization complex formed while undertaking the detection method of claim 2, comprising:
  - a) one strand of a cellular hTRT nucleic acid; and
  - b) one strand of a nucleic acid comprising a-recombinant or synthetic fragment of hTERT, which recombine acid tragment of hTERT comprises at least 40 contiguous nucleotides of <u>consisting</u> separately of 25 or more consecutive nucleotides of the hTERT encoding region of SEQ ID NC:224 or its complement.
- (Withdrawn) The hybridization complex of claim 28, wherein the hTRT nucleic acid is an hTRT mRNA.
- (Withdrawn) The hybridization complex of claim 28, wherein the hTRT nucleic acid is an hTRT cDNA
- (Withdrawn) The hybridization complex of claim 28, wherein the fragment comprises at-least 20 contiguous or more consegutive nucleotides of SEQ ID NO:224 or its complement.

- (Withdrawn) (Currently Amended) The hybridization complex of claim 28, wherein the fragment comprises 10-100-contigueue 100 or more consecutive nucleotides of SEQ ID NO:224 or its complement.
- (Withdrawn) The hybridization complex of claim 28, wherein said hybridization complex is a DNA:DNA complex.
- (Withdrawn) The hybridization complex of claim 28, wherein said hybridization complex is a DNA:RNA complex.
- 35. (New) The method of claim 1, wherein a) comprises combining the sample with the probe at 5°C to 25°C below T<sub>m</sub> in aqueous solution at 1 M NaCl.
- 36. (New) The method of claim 1, wherein the hTRT nucleic acid is hTRT mRNA or cDNA.
- (New) The method of claim 1, wherein the probe comprises a sequence identical or complementary to 100 or more consecutive nucleotides from the hTHT encoding region of SEQ IO NO:224.
- (New) The method of claim 1, wherein the probe comprises a sequence not contained in SEQ. ID NO:62.
- 39. (New) The method of claim 1, wherein the sample has been taken from a patient, and the method further comprises determining or assessing a tumor in the patient according to whether a nucleic acid encoding hTRT or an hTRT fragment is detected.
- 40. (New) The method of claim 2, wherein the sample has been taken from a patient, and the method further comprises determining or assessing a tumor in the patient according to whother said nucleic acid hybrid is detected.
- (New) The method of claim 13, wherein a) comprises combining the sample with the primer at 5°C to 25°C below T<sub>m</sub> in aqueous solution at 1 M NaCl.
- 42. (New) The method of claim 13, wherein the hTRT nucleic acid is mRNA or cDNA.

- 43. (New) The method of claim 13, wherein the primer comprises a sequence identical or complementary to 90 or more consecutive nucleotides from the hTRT encoding region of SEQ ID NO:224.
- (New) The method of claim 13, wherein the primer comprises a sequence not contained in SEQ. ID NO:82.
- 45. (New) The method of claim 13, wherein the sample has been taken from a patient, and the method comprises determining or assessing a tumor in the patient according to whether a nucleic acid encoding hTRT or an hTRT fragment is detected.
- 46. (New) The method of claim 14, wherein the sample has been taken from a patient, and the method comprises determining or assessing a tumor in the patient according to whether said amplification product is formed.